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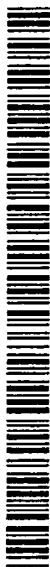


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(54) Title: USE OF PRAMIPEXOLE AS A TREATMENT FOR COCAINE CRAVING

(57) Abstract: Disclosed herein are methods for reducing stimulant dependency or craving, involving administration of a therapeutically-effective amount of a dopamine agonist, such as pramipexole.

USE OF PRAMIPEXOLE AS A TREATMENT FOR COCAINE  
CRAVING

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Background of the Invention

This invention relates to methods for the treatment of cocaine craving.

Cocaine is a highly addictive psychostimulant that causes sensations of euphoria and craving, resulting in physiological as well as psychological damage. Although cocaine use leads to a multitude of physiological complications, its primary target of action is the central nervous system. Cocaine withdrawal following abstinence causes, among other symptoms, an intense craving for the abused drug, which in turn frequently results in the relapse into renewed drug use. Epidemiological studies point to a high incidence of multiple substance abuse among cocaine users, a finding that has significant societal and medical repercussions.

10

To date, approved pharmacotherapies for cocaine abuse and dependence have proven scarce despite the acute need for such therapies.

20

Summary of the Invention

In general, the invention features methods for treating stimulant dependencies, such as cocaine craving, by administering a therapeutically-effective amount of a dopamine agonist, for example, pramipexole.

In one aspect, the invention provides a method of treating a patient (for example, a human) with a stimulant dependency by administering a

25

therapeutically-effective amount of pramipexole to the patient. In preferred embodiments of this aspect, the stimulant dependency is a stimulant craving and the stimulant is cocaine.

5 In a related aspect, the invention provides a method of treating a human diagnosed with cocaine craving by administering a therapeutically-effective amount of pramipexole to the human.

In preferred embodiments of both of the above aspects of the invention, the method further includes administering a therapeutically-effective amount of an antidepressant or an anticonvulsant, for example,  
10 lamotrigine.

By "treating" is meant the medical management of a patient with the intent that a cure, amelioration, or prevention of a dependency or a relapse or associated disease, pathological condition, or disorder will result. This term includes active treatment, that is, treatment directed specifically  
15 toward improvement of the dependency or associated cure of a disease, pathological condition, or disorder, and also includes causal treatment, that is, treatment directed toward removal of the cause of the dependency or associated disease, pathological condition, or disorder. In addition, this term includes palliative treatment, that is, treatment designed for the relief  
20 of symptoms rather than the curing of the dependency, disease, pathological condition, or disorder; preventive treatment, that is, treatment directed to prevention of the dependency or associated disease, pathological condition, or disorder; and supportive treatment, that is, treatment employed to supplement another specific therapy directed toward the improvement of the  
25 dependency or associated disease, pathological condition, or disorder. The term "treating" also includes symptomatic treatment, that is, treatment directed toward constitutional symptoms of the dependency or an associated disease, pathological condition, or disorder.

By "stimulant" is meant any substance that temporarily increases functional activity, and preferably cardiac, respiratory, cerebral, nervous, vascular, motor, or vasomotor functional activity. Preferred stimulants include, without limitation, cocaine, amphetamines, methamphetamine, and  
5 methylphenidate.

By "therapeutically-effective amount" is meant an amount of a pramipexole compound sufficient to produce a healing, curative, or ameliorative effect either in the treatment of a stimulant dependency or in the symptoms of a stimulant dependency, for example, cocaine craving.

10 By "dependency" is meant any form of behavior that indicates an altered or reduced ability to make decisions resulting, at least in part, from the use of stimulants. Representative forms of dependency behavior may take the form of antisocial, inappropriate, or illegal behavior and include those behaviors directed at the desire, planning, acquiring, and use of  
15 stimulants. This term also includes the psychic craving for a drug that may or may not be accompanied by a physiological dependency, as well as a state in which there is a compulsion to take a drug, either continuously or periodically, in order to experience its psychic effects or to avoid the discomfort of its absence. Forms of "dependency" include habituation, that  
20 is, an emotional or psychological dependence on a compound to obtain relief from tension and emotional discomfort, as well as physical or physiological dependence, that is, use of a compound to prevent withdrawal symptoms.

By "craving" is meant a behavior that reflects a consuming desire,  
25 longing, or yearning for a stimulant. This term may refer to aspects of behaviors that are components of a dependency.

The present invention provides a number of advantages.  
Importantly, it provides one of the first therapeutics for the treatment of

stimulant cravings (such as cocaine craving). In addition, the pramipexole utilized herein is non-toxic, is pharmacokinetically understood, and is known to be well tolerated by humans, as is evidenced by its approval for the treatment of Parkinson's Disease.

5                                    Brief Description of the Drawings

Figure 1 is a schematic illustration of the molecular structure of pramipexole, marketed as Mirapex in the United States.

Detailed Description of the Invention

10            The invention described herein features methods involving the administration of pramipexole (or other dopamine-D3/D2 agonists) for the treatment of stimulant dependency, and preferably for the treatment of cocaine craving and its symptoms, as well as cocaine dependency and associated self-destructive behaviors.

15            Described below is an example of the successful use of pramipexole for the treatment of cocaine craving and related symptoms. This example is provided for the purpose of illustrating the invention, and should not be construed as limiting.

Treatment of Cocaine Craving Using Pramipexole

20            Mr. A, a 34 year-old single, successful business man, was referred for evaluation of possible bipolar disorder. Currently depressed, he had in the previous year brought financial ruin on himself by a pattern of cocaine freebasing and sexual and other extravagance that absorbed nearly one million dollars.

25            Along with current major depression, persisting cocaine craving but rare use, and a question of past primary or secondary (to substance abuse)

mania, he manifested an extraordinary movement disorder with constant restlessness and thrashing of his legs, leaving the inner aspects of his knees and thighs bruised and discolored with hematomas in various stages of evolution and resolution.

5           For the restless legs, he had consulted a neurologist who diagnosed “pre-parkinsonism” presumed secondary to neurological damage from cocaine. The disfiguring movements limited his ability to return to and conduct business.

          Previously, he had failed to respond to or tolerate most of the new  
10       generation of antidepressants. Treatment was begun with lamotrigine up to 200 mg with modest improvement in mood. Given his severe restless legs syndrome and persisting depression, pramipexole was added, titrated to 1.5 mg a day in divided doses.

          In response to this treatment, his leg movements quieted  
15       substantially, his mood brightened, and he reported that these were the first days in a year that he awoke without craving cocaine, a benefit sustained for one year on this drug, combined with 75 mg of lamotrigine. During the subsequent year, Mr. A. reported one day of non-compliance when he was out of town without his medication. That night, for the first time, he dreamt  
20       about cocaine and the next day experienced a renewed craving on awakening which resolved when treatment was restored.

          Although he faces an array of financial and business challenges, his mood following treatment is nearly euthymic, his leg movements at worst resemble mild restlessness, and his cocaine craving remains abolished.

25           These dramatic results demonstrate that dopamine agonists, like pramipexole, represent treatments for cocaine craving, and may be particularly useful for patients with comorbid refractory depression.

### Pramipexole and Other Dopamine Agonists

The synthesis of pramipexole is described in U.S. Patent No. 4,886,812 and European Patent 186 087. Pramipexole is a non-ergot derivative which may be used at a range of between about 1.5 mg to 6.0 mg  
5 per day, and is preferably administered between about 1.5 mg and 4.5 mg per day. Higher dosages may be used with the concomitant risk of potential side effects.

Other formulations for treatment or prevention of stimulant dependency or craving, such as cocaine craving, as described herein, may  
10 take the form of a dopamine agonist compound that may be combined with a pharmaceutically-acceptable diluent, carrier, stabilizer, or excipient. Conventional pharmaceutical practice is employed to provide suitable formulations or compositions to administer such compositions to patients. Oral administration is preferred, but any other appropriate route of  
15 administration may be employed, for example, parenteral, intravenous, subcutaneous, intramuscular, intracranial, intraorbital, ophthalmic, intraventricular, intracapsular, intraspinal, intracisternal, intraperitoneal, intranasal, or aerosol administration. Therapeutic formulations may be in the form of liquid solutions or suspensions (as, for example, for intravenous  
20 administration); for oral administration, formulations may be in the form of liquids, tablets or capsules; and for intranasal formulations, in the form of powders, nasal drops, or aerosols.

Methods well known in the art for making formulations are described, for example, in "Remington: The Science and Practice of  
25 Pharmacy" (19th ed.) ed. A.R. Gennaro AR., 1995, Mack Publishing Company, Easton, PA. Formulations for parenteral administration may, for

example, contain excipients, sterile water, saline, polyalkylene glycols such as polyethylene glycol, oils of vegetable origin, or hydrogenated naphthalenes.

If desired, slow release or extended release delivery systems may be utilized. Biocompatible, biodegradable lactide polymer, lactide/glycolide copolymer, or polyoxyethylene-polyoxypropylene copolymers may be used to control the release of the compounds. Other potentially useful parenteral delivery systems include ethylene-vinyl acetate copolymer particles, osmotic pumps, implantable infusion systems, and liposomes. Formulations for inhalation may contain excipients, for example, lactose, or may be aqueous solutions containing, for example, polyoxyethylene-9-lauryl ether, glycocholate and deoxycholate, or may be oily solutions for administration in the form of nasal drops, or as a gel.

In general, a dopamine agonist for use in the methods of the invention is administered at a dosage appropriate to the effect to be achieved and is typically administered in unit dosage form. As noted above, the preferred route of administration for most indications is oral.

An effective quantity of a dopamine agonist-containing compound of the invention is employed to treat the stimulant dependency or craving, for example, cocaine craving as described herein. The exact dosage of the compound may be dependent, for example, upon the age and weight of the recipient, the route of administration, and the severity and nature of the symptoms to be treated. In general, the dosage selected should be sufficient to prevent, ameliorate, or treat the condition, or one or more symptoms thereof, without producing significant toxic or undesirable side effects.



### Combination with Other Therapeutics

One particular source of pramipexole is Pharmacia & Upjohn, Inc. which markets Mirapex (Pramipexole Dihydrochloride) tablets which have the molecular structure shown in Figure 1. Examples of other dopamine  
5 agonists include, but are not limited to, amantadine, bromocriptine, cabergoline, lisuride, pergolide, ropinirole, quinpirole, or quinelorane. Pramipexole, or any other dopamine agonist, may be administered as a monotherapy, or in combination with other compounds, for the treatment of multiple substance abuse or other physiological or psychological conditions.

10 In one particular example, the dopamine agonist (e.g. pramipexole) may be administered in combination with an antidepressant, anticonvulsant, antianxiety, antimanic, antipsychotic, antiobsessional, sedative-hypnotic, or stimulant medication. Examples of these medications include, but are not limited to, the antianxiety medications alprazolam, buspirone hydrochloride,  
15 chlordiazepoxide, chlordiazepoxide hydrochloride, clorazepate dipotassium, desipramine hydrochloride, diazepam, halazepam, hydroxyzine hydrochloride, hydroxyzine pamoate, lorazepam, meprobamate, oxazepam, prazepam, prochlorperazine maleate, prochlorperazine, prochlorperazine edisylate, and trimipramine maleate; the anticonvulsants amobarbital,  
20 amobarbital sodium, carbamazepine, chlordiazepoxide, chlordiazepoxide hydrochloride, clorazepate dipotassium, diazepam, divalproex sodium, ethosuximide, ethotoin, gabapentin, lamotrigine, magnesium sulfate, mephentoin, mephobarbital, methsuximide, paramethadione, pentobarbital sodium, phenacemide, phenobarbital, phenobarbital sodium, phenisuximide,  
25 phenytoin, phenytoin sodium, primidone, secobarbital sodium, trimethadione, valproic acid, and clonazepam; the antidepressants amitriptyline hydrochloride, amoxapine, bupropion hydrochloride, clomipramine hydrochloride, desipramine hydrochloride, doxepin

hydrochloride, fluoxetine, fluvoxamine, imipramine hydrochloride,  
imipramine pamoate, isocarboxazid, lamotrigine, maprotiline  
hydrochloride, nortriptyline hydrochloride, paroxetine hydrochloride,  
phenelzine sulfate, protriptyline hydrochloride, sertraline hydrochloride,  
5    tranylcypromine sulfate, trazodone hydrochloride, trimipramine maleate,  
and venlafaxine hydrochloride; the antimanic medications lithium carbonate  
and lithium citrate; the antiobsessional medications fluvoxamine, and  
clomipramine hydrochloride; the antipsychotic medications acetophenazine  
maleate, chlorpromazine hydrochloride, chlorprothixene, chlorprothixene  
10   hydrochloride, clozapine, fluphenazine decanoate, fluphenazine enathrate,  
fluphenazine hydrochloride, haloperidol decanoate, haloperidol, haloperidol  
lactate, lithium carbonate, lithium citrate, loxapine hydrochloride, loxapine  
succinate, mesoridazine besylate, molindone hydrochloride, perphenazine,  
pimozide, prochlorperazine maleate, prochlorperazine, prochlorperazine  
15   edisylate, promazine hydrochloride, risperidone, thioridazine, thioridazine  
hydrochloride, thiothixene, thiothixene hydrochloride, and trifluoperzine  
hydrochloride; the sedative-hypnotic medications amobarbital, amobarbital  
sodium, aprobarbital, butabarbital, chloral hydrate, chlordiazepoxide,  
chlordiazepoxide hydrochloride, clorazepate dipotassium, diazepam,  
20   diphenhydramine, estazolam, ethchlorvynol, flurazepam hydrochloride,  
glutethimide, hydroxyzine hydrochloride, hydroxyzine pamoate, lorazepam,  
methotrimeprazine hydrochloride, midazolam hydrochloride, non  
prescription, oxazepam, pentobarbital sodium, phenobarbital, phenobarbital  
sodium, quazepam, secobarbital sodium, temazepam, triazolam, and  
25   zolpidem tartrate; and the stimulants dextroamphetamine sulfate,  
methamphetamine hydrochloride, methylphenidate hydrochloride and,  
pemoline.

Other Embodiments

All publications and patent applications mentioned in this specification are herein incorporated by reference to the same extent as if each independent publication or patent application was specifically and  
5 individually indicated to be incorporated by reference.

While the invention has been described in connection with specific embodiments thereof, it will be understood that it is capable of further modifications and this application is intended to cover any variations, uses, or adaptations of the invention following, in general, the principles of the  
10 invention and including such departures from the present disclosure that come within known or customary practice within the art to which the invention pertains and may be applied to the essential features hereinbefore set forth, and follows in the scope of the appended claims.

Other embodiments are within the claims.

15 What is claimed is:

### Claims

1. A method of treating a human with a stimulant dependency, said method comprising administering to said human a therapeutically-effective amount of pramipexole.
- 5           2. The method of claim 1, wherein said stimulant dependency involves a stimulant craving.
3. The method of claim 1, wherein said stimulant is cocaine.
4. A method of treating a cocaine craving in a human, said method comprising administering to said human a therapeutically-effective amount  
10   of pramipexole.
5. The method of claim 1 or 4, wherein said method further comprises administering to said human a therapeutically-effective amount of an antidepressant.
6. The method of claim 1 or 4, wherein said method further  
15   comprises administering to said human a therapeutically-effective amount of an anticonvulsant.
7. The method of claim 6, wherein the anticonvulsant is lamotrigine.

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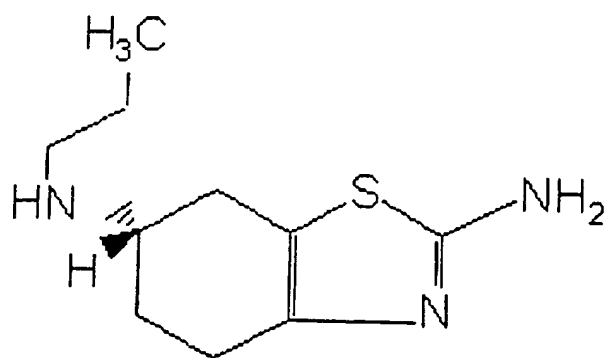


FIGURE 1

## INTERNATIONAL SEARCH REPORT

International application No.  
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<b>A. CLASSIFICATION OF SUBJECT MATTER</b> IPC(7) : A01N 43/78 US CL : 514/367 According to International Patent Classification (IPC) or to both national classification and IPC		
<b>B. FIELDS SEARCHED</b> Minimum documentation searched (classification system followed by classification symbols) U.S. : 514/367  Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched NONE  Electronic data base consulted during the international search (name of data base and, where practicable, search terms used) Please See Extra Sheet.		
<b>C. DOCUMENTS CONSIDERED TO BE RELEVANT</b>		
Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	Database Medline on STN, Department of Neuropharmacology, Scripps Research Institute, (La Jolla, CA), No. 97387623, CAINE, S. B., "D3 Receptor Test in vivo Predicts Decreased Cocaine Self-Administration in Rats", abstract, Neuroreport, 8(9-10), pp. 2373-2377, 07 July 1997.	1-4
<input type="checkbox"/> Further documents are listed in the continuation of Box C. <input type="checkbox"/> See patent family annex.		
* "A" "E" "L" "O" "P"	Special categories of cited documents: document defining the general state of the art which is not considered to be of particular relevance earlier document published on or after the international filing date document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified) document referring to an oral disclosure, use, exhibition or other means document published prior to the international filing date but later than the priority date claimed	*T* "X" "Y" "&*" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art document member of the same patent family
Date of the actual completion of the international search 27 NOVEMBER 2000		Date of mailing of the international search report 04 JAN 2001
Name and mailing address of the ISA/US Commissioner of Patents and Trademarks Box PCT Washington, D.C. 20231 Facsimile No. (703) 305-3230		Authorized officer <i>Dwight C. Jones</i> DWAYNE C. JONES Telephone No. (703) 308-1235

## INTERNATIONAL SEARCH REPORT

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### B. FIELDS SEARCHED

Electronic data bases consulted (Name of data base and where practicable terms used):

REGISTRY, MEDLINE, EMBASE, WPIDS, DRUGU, BIOSIS, CAPLUS structure search with the following terms  
pramipexole?, drugs of abuse, drug dependence, mirapex, snd919 or snd 919, stimulant#(3a)(depend? or additct?),  
lamotrigine.